

(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 0 463 144 B1

(12)

EUROPEAN PATENT SPECIFICATION

(45) Date of publication and mention
of the grant of the patent:
05.02.1997 Bulletin 1997/06

(21) Application number: **91902535.3**

(22) Date of filing: **12.12.1990**

(51) Int Cl.⁶: **C12Q 1/68, G01N 33/543,
C08F 257/02, C08F 285/00,
H01F 1/00, B01J 8/00,
B03C 1/00**

(86) International application number:
PCT/US90/07369

(87) International publication number:
WO 91/09141 (27.06.1991 Gazette 1991/14)

(54) **MAGNETICALLY RESPONSIVE FLUORESCENT POLYMER PARTICLES AND APPLICATION THEREOF**

MAGNETISCHE UND FLUORESZIERENDE POLYMERTeilchen UND IHRE ANWENDUNG
PARTICULES POLYMERES FLUORESCENTES A SENSIBILITE MAGNETIQUE ET LEUR APPLICATION

(84) Designated Contracting States:
AT BE CH DE DK ES FR GB GR IT LI LU NL SE

(30) Priority: **14.12.1989 US 451274**
14.12.1989 US 451483
14.12.1989 US 451494
14.12.1989 US 452099

(43) Date of publication of application:
02.01.1992 Bulletin 1992/01

(73) Proprietor: **DADE INTERNATIONAL INC.**
Deerfield, Illinois 60015-0778 (US)

(72) Inventors:
• **WANG, Chao-Huei, J.**
Gurnee, IL 60031 (US)

• **SHAH, Dinesh, O.**
Vernon Hills, IL 60061 (US)

(74) Representative: **MacGregor, Gordon et al**
ERIC POTTER CLARKSON
St. Mary's Court
St. Mary's Gate
Nottingham, NG1 1LE (GB)

(56) References cited:
WO-A-89/04373 **DE-A- 3 224 484**
US-A- 4 115 535 **US-A- 4 206 094**
US-A- 4 369 226 **US-A- 4 563 419**
US-A- 4 707 523 **US-A- 4 891 324**

Exempted from 395, 688

Note: Within nine months from the publication of the mention of the grant of the European patent, any person may give notice to the European Patent Office of opposition to the European patent granted. Notice of opposition shall be filed in a written reasoned statement. It shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

EP 0 463 144 B1

D scription

This invention relates to processes using magnetically responsive fluorescent polymer particles.

5 Background of the Invention

Many biological techniques, such as immunoassays, affinity purification etc., require the separation of bound from free fractions. Magnetic particles have been used to facilitate the desired separation.

10 Magnetic particles have been formed from a variety of particulate and magnetic matter, using a variety of processes, having different characteristics. For example, US-A 4,582,622, discloses a magnetic particle comprised of gelatin, water-soluble polysaccharide, sodium phosphate and ferromagnetic substances; US-A 4,628,037 and US-A-4,554,088 disclose magnetic particles comprised of a magnetic metal oxide core surrounded by a coat of polymeric silane; US-A-4,452,773 discloses discrete colloidal sized particles having a core of ferromagnetic iron oxide (Fe_3O_4) which is coated with a water-soluble polysaccharide or a derivative thereof having functional groups; and US-A-4,297,337 dis-

15 closes magnetic glass- or crystal-containing material as a particulate carrier. WO-A-89/04373 discloses monodispersed magnetic particles of uniform size distribution and magnetic content and having a polymer inner core and a coating of magnetically responsive metal oxide polymerised with a monomer. The coating is able to adsorb to the inner core.

20 The document also discloses a process for using such particles for the determination of the presence or concentration of an analyte and for determining the presence or concentration of specific nucleic acid sequences in nucleic acid target molecules.

The present invention provides such processes in which the particles are fluorescent and are used to monitor the number of particles present during the processes. The processes are as defined in Claims 1 and 2.

US-A-4206094 and US-A-4707523 disclose magnetic particles which can be labelled with fluorescent dye.

25 The magnetic particles may be prepared by first producing a magnetically responsive metal oxide, hereinafter referred to as metal oxide, with average size of about 1 micron or less and then coating a fluorescent polymeric core particle of average size 1 to 100 μm with a layer of polymer containing metal oxide. The surface of these fluorescent magnetic particles can be coated further with another layer of polymer or functionalized polymer to provide the desired surface characteristics.

30 The spectral characteristics of these fluorescent magnetic particles can be varied by using core particles incorporated with various fluorescent dyes, either a single fluorescent dye or a combination of several fluorescent dyes. Alternatively, the fluorescent magnetic particles can be prepared by incorporating various fluorescent dyes, either a single fluorescent dye or a combination of several fluorescent dyes, which are soluble in the monomer and can withstand the polymerization condition, in the presence of nonfluorescent polymeric core particles, metal oxide, and monomer.

35 The fluorescent material may be selected from NileTM red, CoumarinTM6, CoumarinTM4, RhodamineTMB, NileTMblue, oxazine 725, oxazine 750, or a mixture of one of these fluorescent materials with one or more other of these materials.

40 The fluorescent magnetic particles are monodispersed in size with rough surface and have a magnetic metal oxide content of from about 5% to 50%, preferably from 10% to 25%. The fluorescence intensity of these fluorescent magnetic particles can be adjusted by varying the magnetic metal oxide content to vary the shading due to the metal oxide and/or varying the amount of fluorescent dye incorporated into the fluorescent polymeric core particles.

45 Particles with these characteristics have been found to be useful in immunoassays and a wide variety of biomedical applications. These fluorescent magnetic particles can be used for passive or covalent coupling of biological material such as antigens, antibodies, enzymes or DNA/RNA and used as solid phase for various types of immunoassays, DNA/RNA hybridization assays, affinity purification, cell separation, phagocytosis, and other biomedical applications. These fluorescent magnetic particles with or without coupling to biological material can be incorporated at various ratios to nonmagnetic particles for various assays to serve as markers for making sure that the correct number of particles are being delivered into the well and for checking the particle loss during the assay.

Both crosslinked and noncrosslinked magnetically responsive fluorescent polymer particles can be used.

50 Monodispersed magnetically responsive fluorescent polymer particles can be used.

Detailed Description of the Invention

55 The fluorescent magnetic particles may be prepared by first producing metal oxide with average size of about 1 micron or less. The metal oxide is produced by heating and precipitating a mixture of divalent and trivalent metal salt, preferably a mixture of ferrous and ferric sulfate or chloride with sodium hydroxide solution. The molar ratio of divalent to trivalent metal salt can be varied from 0.5 to 2.0, preferably 0.5 to 1.0, to obtain the desirable size and magnetic characteristics of metal oxide. It is observed that the molar ratio of divalent to trivalent metal salt affects the size of the

metal oxide: the smaller the molar ratio of divalent to trivalent metal salt, the smaller the size of metal oxide. The molar ratio of divalent to trivalent metal salt also affects the color of the resulting magnetic particles: the smaller the molar ratio, the lighter the brownish color of the resulting magnetic particles. Preferably, the metal oxide is either superparamagnetic or paramagnetic although ferromagnetic metal oxide can also be used, provided centrifugation instead of magnetic separation is used during the clean up.

Other divalent transition metal salts such as manganese, magnesium, cobalt, nickel, zinc, and copper salts may be substituted for ferrous salt.

After the metal oxide has been precipitated, it is washed several times with centrifugation at 250xg until the supernatant is neutral in pH. The metal oxide is resuspended in deionized water and mechanically stirred at high speed to break down the aggregate of metal oxide crystals. Further centrifugation at 250xg will not pellet all of the metal oxide. The supernatant which contains smaller size metal oxide crystals is collected and the pellet is resuspended in deionized water. This process is repeated for at least three times or until most of metal oxide can no longer be pelleted at 250xg. The metal oxide obtained this way usually has size less than 2.0 micron. Low speed centrifugation at 100xg to remove larger crystals will reduce the size to less than 0.8 micron.

The metal oxide with average size of 1.0 micron or less is mixed with monomer and coated onto the fluorescent polymeric core particles, preferably polystyrene particles, with size of 1 to 100 microns in the presence of initiator. Addition of a small quantity of emulsifier will help prevent the particles from agglomerating. The fluorescent magnetic particles are then coated with a protective layer of polymer, preferably polystyrene, to prevent the metal oxide from falling off. If functionalized fluorescent magnetic particles are desired, the magnetic particles can be coated further with another layer of functionalized polymer to provide functional groups such as carboxyl, amino or hydroxyl for covalent coupling of biological material.

The fluorescent magnetic particles can be illustrated in Figure I, where 1 represents the fluorescent core particle, 2 represents the metal oxide/polymer coating, 3 represents the protective polymer coating and 4 represents the functionalized polymer coating. Figure II shows the transmission electron micrograph of 0.08 to 0.1 micron slice of a magnetic particle-prepared as described above. Figure III shows a scanning electron micrograph of 6.8 micron magnetic particles. Figure III a is at 1000x and Figure III b is at 5000x magnification.

The fluorescent polymeric core particles useful in this invention may be of any polymer which can be obtained as a dispersion of small particles and which can absorb a monomer thereby causing the metal oxide and monomer mixture to coat onto the surface of the core particles. The core particles may be of any size and shape, preferably of 1 to 100 microns in size and spherical in shape. When monodispersed core particles are used the resulting magnetic particles will also be monodispersed in size. The core particles may be obtained by emulsion polymerization, suspension polymerization or other means of polymerization with or without a crosslinking agent such as divinyl benzene or the like. Among the monomers which can be used to prepare core particles are styrene, methyl methacrylate, vinyltoluene and the like. A mixture of the monomers can also be used. The fluorescent core particles may be obtained by incorporating the fluorescent dye in the core particles, using various techniques known to those skilled in the art. The monomer used for magnetic metal oxide coating or protective coating may or may not be the same type as the fluorescent core particles. The weight ratio of monomer used for metal oxide coating to fluorescent core particles may be from 0.1 to 12, preferably from 0.2 to 6, depending upon the thickness of metal oxide/polymer layer desired. When the metal oxide prepared from a mixture of ferrous and ferric salts is used for coating it is preferred to use a monomer to fluorescent core particle weight ratio of about 0.1 to 0.5. However when the metal oxide prepared from a mixture of manganese (II) and ferric salts is used for coating the weight ratio of monomer to core particles may be from 0.1 to 12. As a result when crosslinked fluorescent magnetic particles which are inert to common organic solvent are desired, it is preferred to use the metal oxide prepared from a mixture of manganese (II) and ferric salts with monomer containing 2% to 10%, preferably 8% to 10% by weight of crosslinking agent and a monomer to core particle weight ratio of 3 to 12, preferably 4 to 6 and a monomer to fluorescent dye weight ratio of 0.1 to 10. When lower monomer to core particle weight ratio (i.e. 0.1 to 0.5) is used during the metal oxide/polymer coating it is preferred to overcoat the resulting fluorescent magnetic particles with a protective layer of polymer coating to further adhere the metal oxide to the surface of the fluorescent magnetic particles. However, when higher monomer to core particle ratio (i.e. 3 to 12) is used no protective polymer coating is necessary. The polymerization temperature may be from 50°C to 90°C, preferably 55°C to 65°C. The polymerization initiator may either be water soluble such as potassium persulfate and the like or water insoluble such as benzoyl peroxide and the like. Other means of polymerization initiation such as radiation, ionization or the like may also be used. It is found unexpectedly that fluorescent magnetic particles can be produced without using any emulsifier when the metal oxide prepared from a mixture of manganese (II) and ferric salts is used for coating. However, a small amount of emulsifier such as sodium dodecylsulfate, Aerosol™ 22, Tween™ 20 or Nonidet™ P-40 (NP 40) is found to be useful in preventing the particles from extensive aggregation during the metal oxide/polymer coating when the metal oxide prepared from a mixture of ferrous and ferric salts is used for coating. Other emulsifiers with the same capability may also be used. The magnetic metal oxide content can be varied from 5% to 50%, preferably from 10% to 25% by using different amounts of metal oxide during the metal oxide/polymer coating. Multiple metal oxide/polymer coatings can

also be employed to increase the metal oxide content. Other ingredients commonly used in polymerization may also be added as long as magnetic particles with desirable characteristics can be obtained. The ingredients for metal oxide/polymer coating may be added all at once at the beginning of metal oxide/polymer coating process or added stepwise. When the metal oxide prepared from a mixture of ferrous and ferric salt is used, it is preferred to add the ingredients stepwise. The ingredients may be mixed by mechanical stirring, tumbling or other means of agitation under vacuum or inert gas such as argon. The functional groups can be incorporated onto the surface of the fluorescent magnetic particles by either using a mixture of monomer and functionalized monomer during the metal oxide/polymer coating or overcoating the magnetic particles with a thin layer of functionalized monomer at the end. The functionalized monomer used may be selected from one or a mixture of the following: 2-hydroxyethyl methacrylate, 2-aminoethyl methacrylate, trimethylammoniummethyl methacrylate methosulfate, dimethylaminoethyl methacrylate, methacrylic acid, undecylenic acid, methyl propene sulfonic acid, undecylenyl alcohol, oleyl amine, glycidyl methacrylate, acrolein, glutaraldehyde and the like. The magnetic fluorescent particles can also be overcoated with a layer of different polymer than the one used for metal oxide/polymer coating or protective coating to take up the surface characteristics of that polymer.

Applications of Fluorescent Magnetic Particles

The uses of a wide variety of fluorescent magnetic particles as solid phase for various applications such as fluorescence immunoassays, radioimmunoassays, enzyme immunoassays, cell separations, enzyme immobilizations and affinity purifications have been reviewed in literature as exemplified by the following articles: Hirschbein et al, Chemical Technology, March 1982, 172-179 (1982); Pourfarzaneh, The Ligand Quarterly, 5(1): 41-47 (1982); Halling and Dunnill, Enzyme Microbe Technology, 2: 2-10 (1980); Mosbach and Anderson, Nature, 270: 259-261 (1977); Guesdon et al, J. Allergy Clinical immunology, 61(1), 23-27 (1978). Some applications have also been disclosed in the U.S. Patent Nos. 4,152,210 and 4,343,901 for enzyme immobilizations; U.S. Patent Nos. 3,970,518, 4,230,685, and 4,267,2343 for cell separations; U.S. Patent Nos. 4,554,088, 4,628,037, and 3,933,997 for immunoassays.

Some magnetic particles may be useful in one application, but not in another application. For example, the magnetic particles disclosed in U.S. Patent No. 4,554,088 and 4,628,037, which comprise a superparamagnetic metal oxide core generally surrounded by a coat of polymeric silane, may be useful in immunoassay and affinity purification, due to the large surface area and slower settling rate, but are not suitable in cell separation application such as bone marrow purging. Due to the small size of the magnetic particles, disclosed in these two patents, it is very difficult to remove all of the magnetic particles from the cell suspension effectively. Moreover, the nonspecific binding of smaller magnetic particles to normal cells would be much higher. In using magnetic particles for bone marrow purging, the magnetic particles are coated with antibody, such as sheep anti-mouse IgG, and the bone marrow is treated with a mixture of several monoclonal antibodies against the cancer cell surface antigens. The magnetic particles will bind only to the cancer cells and cause them to be separated from normal cells by passing them through a strong magnetic field. The cleansed cells are then put back into the patient.

The magnetic particles used in the present invention can be optimized in terms of size, surface area, metal oxide content and surface characteristics for a wide variety of biomedical applications. The magnetic particles can be used as solid phase for enzyme immunoassay, fluorescence immunoassay, radioimmunoassay, DNA/RNA hybridization assay, and other diagnostic applications. Immunoassays can be performed by using various configurations such as sandwich assays and competitive binding assays etc., which are obvious to those skilled in the art. The DNA/RNA hybridization can also be performed by using various configurations such as solid phase hybridization or liquid phase hybridization. In solid phase hybridization configuration a DNA or RNA probe (catcher probe) is immobilized on the magnetic particle first. The immobilized catcher probe is then used to hybridize with complimentary strand of DNA from the sample (sample DNA). Finally another probe (signal probe) which is labeled with fluorescent, radioactive or enzyme tracer and capable of hybridizing with another part of the sample DNA is used for signal generation. In liquid phase hybridization configuration the catcher probe and signal probe are allowed to hybridize with the sample DNA in the liquid phase first and are then immobilized to the magnetic particles.

Alternatively, the signal probe can also be labelled with one or several biotin groups and the signal is detected by binding the biotin groups with avidin labelled fluorescent, radioactive or enzymatic tracer to enhance the sensitivity of the assay.

The immunoassays and DNA/RNA hybridization assays can be used to measure a wide variety of compounds such as drugs, hormones, antibodies, peptides, DNA, RNA, nucleotides, viral antigens, and carbohydrates in biological samples.

The magnetic particles can also be used for affinity purification, cell separation, enzyme immobilization and other biomedical applications. In cell separation the magnetic particles are used to either remove unwanted cells (negative selection) or enrich the wanted cells (positive selection) through immunological reactions or nonimmunological reactions. This principle can be used to remove cancer cells from bone marrow (bone marrow purging), purify cell populations through either positive or negative selection for tissue culture and perform various cellular immunoassays etc. In

affinity purification the magnetic particles are used in place of conventional solid phase such as polyacrylamide gels, sepharose gels or other cellulose beads to purify a wide variety of biological materials such as antibodies, antigens, enzymes, inhibitors, cofactors, single stranded DNA, binding proteins, haptens and carbohydrates etc. In another application similar to the affinity purification, the magnetic particles can be used to cross adsorb and remove unwanted protein components from the antisera or clinical samples. In enzyme immobilization the enzyme is immobilized onto the magnetic particles through various means of coupling so as to preserve the enzyme activity and to permit the reuse of immobilized enzyme. The magnetic particles with immobilized enzyme can be used to replace other solid phases such as glass beads, controlled pore glass, silica gels and cellulose beads etc., which are commonly used in immobilized enzyme systems to produce a wide variety of materials such as carbohydrates, amino acids, and proteins, etc.

These fluorescent magnetic particles with or without coupling to biological material can be incorporated at various ratios to nonfluorescent particles for various assays as mentioned in Examples 42 and 43 to serve as markers for making sure that the correct number of particles are being delivered into the well and for checking the particle loss during the assay.

These applications are all facilitated by the ease of separation, fast reaction rate and large surface area common to most of magnetic particles. The following examples are provided to further illustrate the versatility and advantages of this invention.

General Procedures for the Preparation of Metal Oxide

Example 1

In a three-necked round bottom flask equipped with mechanical stirrer, condenser, thermometer, dropping funnel and heating mantle was placed a mixture containing 0.361 mol of ferrous sulfate and 0.369 mol of ferric sulfate ($\text{Fe}^{++}/\text{Fe}^{+++}$ ratio = 1) in 400 ml of deionized water. The mixture was heated to 85 to 90 °C with stirring and added dropwise 850 ml of 6 N sodium hydroxide over a period of 90 minutes. The mixture was stirred at 85 to 90 °C for one more hour and cooled to room temperature. The metal oxide precipitates were centrifuged at 250xg for 10 minutes. The clear supernatant was decanted and the pellet was resuspended in 900 ml of deionized water using mechanical stirrer. This cleaning process was repeated six times or until the supernatant was almost neutral in pH. The supernatant was decanted and resuspended in 200 ml of deionized water. Further centrifugation at 250xg will not pellet all of the metal oxide precipitates. The supernatant which contained smaller size metal oxide crystals was collected and the pellet was resuspended in 200 ml of deionized water. This process was repeated for at least three times or until most of metal oxide can no longer be pelleted at 250xg. The metal oxide obtained this way usually has size less than 2.0 micron. The combined metal oxide suspension was centrifuged at 100xg for 10 minutes. The supernatant was collected to give 800 ml of 8.6 % w/v magnetic metal oxide suspension having the size less than 0.8 microns.

Example 2

Same procedures as described in Example 1 were followed except 0.235 mol of ferrous sulfate, 0.297 mol of ferric sulfate ($\text{Fe}^{++}/\text{Fe}^{+++}$ ratio = 0.79) in 400 ml of deionized water and 480 ml of 6 N sodium hydroxide were used to give 2000 ml of 2.86 % w/v suspension of magnetic metal oxide.

Example 3

Same procedures as described in Example 1 were followed except 0.178 mol of ferrous sulfate, 0.298 mol of ferric sulfate ($\text{Fe}^{++}/\text{Fe}^{+++}$ ratio = 0.59) in 400 ml of deionized water and 520 ml of 6 N sodium hydroxide were used to give 1500 ml of 2.98 % w/v suspension of magnetic metal oxide.

Example 4

Same procedures as described in Example 1 were followed except 0.15 mol of ferrous sulfate, 0.276 mol of ferric sulfate ($\text{Fe}^{++}/\text{Fe}^{+++}$ ratio = 0.54) in 400 ml of deionized water and 520 ml of 6 N sodium hydroxide were used to give 700 ml of 6.88 % w/v suspension of magnetic metal oxide.

Example 5

Same procedures as described in Example 1 were followed except 0.116 mol of manganese sulfate, 0.146 mol of ferric sulfate ($\text{Mn}^{++}/\text{Fe}^{+++}$ ratio = 0.79) in 225 ml of deionized water and 240 ml of 6 N sodium hydroxide were used to give 1700 ml of 1.8 % w/v suspension of magnetic metal oxide.

Preparation of Magnetic ParticlesExample 6

5 A mixture containing 600 ml of deionized water, 6 ml of styrene and 80 ml of 8.6 % w/v magnetic metal oxide prepared as described in Example 1, was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for one hour. To the mixture were added 12 g of potassium persulfate and 850 ml of 5 % w/v, 4.0 micron polystyrene particles. The bottle was resealed, evacuated and rotated for one hour and added 50 ml of 2 % sodium dodecylsulfate. After five more hours 6 ml of styrene and 10 g of potassium persulfate were added to the mixture. The mixture was rotated for another fifteen hours, filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting magnetic particles were resuspended to 1.6 l with deionized water to give a 2.5 % w/v suspension with about 11 % magnetic metal oxide content and 4.3 micron average size.

Example 7

15 The magnetic particles, 1.6 l of 2.5 % w/v, prepared as described in Example 6, were carboxylated by adding 1 g of sodium dodecylsulfate, 10 g of potassium persulfate and a solution containing 0.98 ml of undecylenic acid and 0.02 ml of divinyl benzene in 4 ml of methanol. The mixture was placed in a sealed bottle, evacuated and rotated at about 60 rpm in a 55 °C oven for 5 hours. The resulting carboxyl magnetic particles were separated magnetically and washed several times with deionized water until the supernatant was clear. The carboxyl magnetic particles were resuspended to 680 ml with deionized water to give a 5.8 % w/v suspension with about 11 % magnetic metal oxide content and 4.3 micron average size.

Example 8

25 A mixture containing 600 ml of deionized water, 6 ml of styrene and 80 ml of 8.6 % w/v magnetic metal oxide prepared as described in Example 1, was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for one hour. To the mixture were added 12 g of potassium persulfate and 850 ml of 4.78 % w/v, 6.1 micron polystyrene particles. The bottle was resealed, evacuated, rotated for five hours and added 6 ml of styrene and 10 g of potassium persulfate. The mixture was rotated for another fifteen hours, filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting magnetic particles were resuspended to 1.5 l with deionized water and carboxylated by adding 1 g of sodium dodecylsulfate, 10 g of potassium persulfate and a solution containing 0.98 ml of undecylenic acid and 0.02 ml of divinyl benzene in 4 ml of methanol. The mixture was placed in a sealed bottle, evacuated and rotated at about 60 rpm in a 55 °C oven for 5 hours. The resulting carboxyl magnetic particles were separated magnetically and washed several times with deionized water until the supernatant was clear. The carboxyl magnetic particles were resuspended to 800 ml with deionized water to give a 4.3 % suspension with about 11.6 % magnetic metal oxide content and 6.8 micron average size.

Example 9

45 A mixture containing 600 ml of deionized water, 6 ml of styrene and 60 ml of 8.6 % w/v magnetic metal oxide prepared as described in Example 1, was placed in a three-necked round bottom flask and stirred at 67 °C for one hour under argon. To the mixture were added 12 g of potassium persulfate and 470 ml of 5 % w/v, 2.7 micron polystyrene particles. The mixture was stirred at 67 °C for one hour and added 30 ml of 2 % sodium dodecylsulfate. After stirring at 67 °C under argon for five more hours 6 ml of styrene and 6 g of potassium persulfate were added to the mixture. The mixture was stirred at 67 °C under argon for another fifteen hours, filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting magnetic particles were resuspended to 900 ml with deionized water and carboxylated by adding 0.6 g of sodium dodecylsulfate, 10 g of potassium persulfate and a solution containing 0.598 ml of undecylenic acid and 0.012 ml of divinyl benzene in 2.4 ml of methanol. The mixture was placed in a sealed bottle, evacuated and rotated at about 60 rpm in a 55 °C oven for 5 hours. The resulting carboxyl magnetic particles were separated magnetically and washed several times with deionized water until the supernatant was clear. The carboxyl magnetic particles were resuspended to 500 ml to give a 6.5 % w/v suspension with about 14 % magnetic metal oxide content and 4.0 micron average size.

Example 10

A mixture containing 600 ml of deionized water, 6 ml of styrene and 60 ml of 8.6 % w/v magnetic metal oxide prepared as described in Example 1, was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for one hour. To the mixture were added 12 g of potassium persulfate and 470 ml of 5 % w/v, 2.7 micron polystyrene particles. The bottle was resealed, evacuated and rotated for one hour and added 30 ml of 2 % sodium dodecylsulfate. After five more hours 6 ml of styrene and 10 g of potassium persulfate were added to the mixture. The mixture was rotated for another fifteen hours, filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting magnetic particles were resuspended to 500 ml with deionized water to give a 6.8 % w/v suspension with about 14 % magnetic metal oxide content and 4.0 micron average size.

Example 11

A mixture containing 180 ml of deionized water, 2 ml of styrene and 20 ml of 8.6 % w/v magnetic metal oxide, prepared as described in Example 1, was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for one hour. To the mixture were added 4 g of potassium persulfate and 160 ml of 6.8 % w/v magnetic particles (3.0 micron, 14 % metal oxide content), prepared as described in Example 10. The bottle was resealed, evacuated and rotated for one hour and added 10 ml of 2 % sodium dodecylsulfate. After 5 more hours 2 ml of styrene and 2 g of potassium persulfate were added to the mixture. The mixture was rotated for another fifteen hours, filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting magnetic particles were resuspended to 160 ml with deionized water to give a 7.78 % w/v suspension with about 19 % metal oxide content and 4.2 micron average size.

Example 12

A mixture containing 90 ml of deionized water, 1 ml of styrene and 10 ml of 8.6 % w/v magnetic metal oxide, prepared as described in Example 1, was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for one hour. To the mixture were added 1 g of potassium persulfate and 80 ml of 7.78 % w/v magnetic particles (3.2 micron, 19 % metal oxide content), prepared as described in Example 11. The bottle was resealed, evacuated and rotated for four hour and added 5 ml of 2 % sodium dodecylsulfate. After 5 more hours 1 ml of styrene and 1 g of potassium persulfate were added to the mixture. The mixture was rotated for another fifteen hours, filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting magnetic particles were resuspended to 160 ml with deionized water to give a 4.5 % w/v suspension with about 23 % metal oxide content and 4.5 micron average size.

Example 13

A mixture containing 400 ml of deionized water, 1.92 ml of styrene, 0.08 ml of divinyl benzene, 4 g of potassium persulfate, 20 g of 200 - 400 mesh 4 % divinyl benzene cross linked polystyrene beads and 10 ml of 8.6 % w/v magnetic metal oxide, prepared as described in Example 1, was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for 15 hours. The mixture was allowed to settle and the supernatant was decanted. The resulting magnetic beads were resuspended in 200 ml of deionized water and allowed to settle again. This process was repeated several times until the supernatant was clear. The resulting magnetic beads were resuspended in 200 ml of deionized water and added 0.1 g of sodium dodecyl sulfate, 2.0 g of potassium persulfate, 0.48 ml of styrene, and 0.02 ml of divinyl benzene. The bottle was resealed, evacuated and rotated at about 60 rpm in a 55 °C oven for one hour and added a solution containing 0.098 ml of undecylenic acid and 0.002 ml of divinyl benzene in 0.4 ml of methanol. The mixture was rotated for four more hours and cleaned up by gravitational sedimentation as described previously. The water was removed by filtration and the carboxyl magnetic beads were dried to give 20 g of 200 - 400 mesh carboxyl magnetic beads.

Example 14

A mixture containing 100 ml of deionized water, 0.5 ml of styrene, 2 g of potassium persulfate, 75 ml of 5 % w/v 4.0 micron polystyrene particles and 10 ml of 6.88 % w/v magnetic metal oxide, prepared as described in Example 4, was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for fifteen hours. The mixture was filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting magnetic particles were resuspended to 150 ml with

deionized to give a 2.5 % w/v suspension with about 14 % metal oxide content and 4.3 micron average size.

Example 15

Same procedures as described in Example 14 were followed except 20 ml of 6.88 % w/v magnetic metal oxide, prepared as described in Example 4, was used to give 160 ml of 2.5 % w/v suspension with about 18 % metal oxide content and 4.3 micron average size.

Example 16

A mixture containing 2000 ml of deionized water, 13 ml of styrene and 550 ml of 2.98 % w/v magnetic metal oxide prepared as described in Example 3, was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for one hour. To the mixture were added 20 g of potassium persulfate and 950 ml of 10 % w/v, 3.0 micron polystyrene particles. The bottle was resealed, evacuated and rotated for one hour and added 60 ml of 2 % sodium dodecylsulfate. After five more hours 8 ml of styrene and 10 g of potassium persulfate were added to the mixture. The mixture was rotated for another fifteen hours, filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting magnetic particles were resuspended to 3000 ml with deionized water to give a 3.38% w/v suspension with about 12 % magnetic metal oxide content and 3.2 micron average size.

Example 17

A mixture containing 150 ml of magnetic particles (3.2 micron, 3.38 % w/v with 12 % metal oxide content) prepared as described in Example 16, 2 ml of 1 % NP 40, 0.5 ml of methyl methacrylate or styrene, 1 g of potassium persulfate and 2 ml of functionalized monomer, trimethylammoniummethyl methacrylate methosulfate (40 % aqueous solution), was placed in a sealed bottle. The bottle was rotated at about 60 rpm in a 55 °C oven for four hours. The mixture was filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting magnetic particles were resuspended to 200 ml with deionized water to give a 2.5 % w/v suspension of magnetic particles with trimethylammonium functional groups on the surface.

Example 18

Same procedures as described in Example 17 were followed except 1 ml of functionalized monomer, 2-aminoethyl methacrylate, was used to give 200 ml of 2.5 % w/v suspension of magnetic particles with amino groups on the surface.

Example 19

Same procedures as described in Example 17 were followed except 1 ml of functionalized monomer, 2-hydroxyethyl methacrylate, was used to give 200 ml of 2.5 % w/v suspension of magnetic particles with hydroxyl groups on the surface.

Example 20

Same procedures as described in Example 17 were followed except 1 ml of monomer, 1-vinyl-2-pyrrolidinone, was used to give 200 ml of 2.5 % w/v suspension of magnetic particles with polyvinylpyrrolidinone on the surface.

Example 21

Same procedures as described in Example 17 were followed except 1 g of functionalized monomer, methyl propene sulfonic acid, was used to give 200 ml of 2.5 % w/v suspension of magnetic particles with sulfonic acids groups on the surface.

Example 22

Same procedures as described in Example 17 were followed except 1 ml of functionalized monomer, dimethylaminoethyl methacrylate, was used to give 200 ml of 2.5 % w/v suspension of magnetic particles with dimethylamino groups on the surface.

Example 23

A mixture containing 20 ml of 7.0 % w/v, 2.11 micron polystyrene particles, 100 ml of 1.8 % w/v metal oxide prepared as described in Example 5, 50 ml of deionized water and a solution containing 0.15 g of benzoyl peroxide in 7.5 ml of styrene was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for fifteen hours. The mixture was filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting magnetic particles were resuspended to 200 ml with deionized water to give 5.0 % w/v suspension with about 16.8 % metal oxide content and 3.6 micron average size.

Example 24

A mixture containing 20 ml of 7.0 % w/v, 2.11 micron polystyrene particles, 100 ml of 1.8 % w/v metal oxide prepared as described in Example 5, 50 ml of deionized water and a solution containing 0.15 g of benzoyl peroxide and 0.75 ml of divinyl benzene in 6.75 ml of styrene was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for fifteen hours. The mixture was filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting crosslinked magnetic particles were resuspended to 200 ml with deionized water to give 5.0 % w/v suspension with about 16.8 % metal oxide content and 3.6 micron average size. The crosslinked magnetic particles prepared this way were found to be uniform in size and inert to common organic solvents such as acetone, acetonitrile and dimethyl formamide.

Example 25

A mixture containing 20 ml of 7.0 % w/v, 2.11 micron polystyrene particles, 150 ml of 1.8 % w/v metal oxide prepared as described in Example 5 and a solution containing 0.15 g of benzoyl peroxide, 0.75 ml of divinyl benzene in 6.75 ml of styrene was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for fifteen hours. The mixture was filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting crosslinked magnetic particles were resuspended to 200 ml with deionized water to give 5.4 % w/v suspension with about 23 % metal oxide content and 4.0 micron average size. The crosslinked magnetic particles prepared this way were found to be uniform in size and inert to common organic solvents such as acetone, actonitrile and dimethyl formamide.

Example 26

A mixture containing 15 ml of 9.16 % w/v, 3.2 micron polystyrene particles, 100 ml of 1.8 % w/v metal oxide prepared as described in Example 5, 55 ml of deionized water and a solution containing 0.15 g of benzoyl peroxide and 0.75 ml of divinyl benzene in 6.75 ml of styrene was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for fifteen hours. The mixture was filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting crosslinked magnetic particles were resuspended to 200 ml with deionized water to give 4.7 % w/v suspension with about 16.8 % metal oxide content and 5.5 micron average size. The crosslinked magnetic particles prepared this way were found to be uniform in size and inert to common organic solvents such as acetone, actonitrile and dimethyl formamide.

Example 27

A mixture containing 30 ml of 4.5 % w/v, 4.1 micron polystyrene particles, 100 ml of 1.8 % w/v metal oxide prepared as described in Example 5, 40 ml of deionized water and a solution containing 0.15 g of benzoyl peroxide and 0.75 ml of divinyl benzene in 6.75 ml of styrene was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a 55 °C oven for fifteen hours. The mixture was filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting crosslinked magnetic particles were resuspended to 200 ml with deionized water to give 4.5 % w/v suspension with about 16.9 % metal oxide content and 6.7 micron average size. The crosslinked magnetic particles prepared this way were found to be uniform in size and inert to common organic solvents such as acetone, acetonitrile and dimethyl formamide.

Example 28

A mixture containing 20 ml of 7.0 % w/v, 2.11 micron polystyrene particles, 100 ml of 1.8 % w/v metal oxide prepared as described in Example 5, 50 ml of deionized water and a solution containing 0.15 g of benzoyl peroxide, 0.75 ml of undecylenyl alcohol and 0.75 ml of divinyl benzene in 6 ml of styrene was placed in a sealed bottle. The bottle was

evacuated and rotated at about 60 rpm in a 55 °C oven for fifteen hours. The mixture was filtered through two layers of cheese cloth, separated magnetically and washed several times with deionized water until the supernatant was clear. The resulting crosslinked hydroxyl magnetic particles were filtered and dried to give 9 g of powder with about 16.8 % metal oxide content and 3.9 micron average size. The crosslinked hydroxyl magnetic particles prepared this way were found to be uniform in size and inert to common organic solvents such as acetone, acetonitrile and dimethyl formamide.

Coupling Biological Materials to Magnetic Particle

Example 29

In a 80 ml bottle was placed 30 ml of 4.3 micron, 5.0 % w/v carboxyl magnetic particles prepared as described in Example 7. The particles were separated magnetically and resuspended in 50 ml of phosphate buffer (0.1 M, pH 5.5). To the particle suspension were added 20 mg of bovine serum albumin and 100 mg of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide (EDC). The mixture was rotated end to end at room temperature for two hours and separated magnetically. The particles were washed once with 80 ml of phosphate buffer and resuspended to 75 ml with phosphate buffered saline (0.1 M, pH 7.0) to give a 2.0 % w/v suspension.

To couple bovine serum albumin to magnetic particles by passive adsorption the same procedures were followed except no EDC was used.

Example 30

In a 4 ml vial was placed 1 ml of 4.3 micron, 5.0 % w/v carboxyl magnetic particles prepared as described in Example 7. The particles were separated magnetically and washed once with 2 ml of phosphate buffer (0.1 M, pH 5.5) and resuspended to 2 ml with the same buffer. To the particles suspension were added 140 µl of 1.4 mg/ml Goat (Gt) anti Mouse (Ms) IgG and 10 mg of 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide. The vial was rotated end to end at room temperature for two hours. The particles were separated magnetically, washed once with 2 ml of phosphate buffer and resuspended to 2 ml with phosphate buffered saline (0.1 M, pH 7.0) to give a 2.5 % w/v Gt anti Ms IgG coated magnetic particles. Other kind of antibody either monoclonal or polyclonal could also be coupled to carboxyl magnetic particles by using the same procedures.

To couple Gt anti Ms IgG or other kind of antibody to the magnetic particles by passive adsorption the same procedures were followed except no EDC was used.

Example 31

In a 4 ml vial was placed a 2.5 ml of bovine serum albumin coated magnetic particles (4.3 micron, 2 % w/v) prepared as described in Example 29. The particles were separated magnetically and resuspended to 2 ml with phosphate buffer (0.1 M, pH 5.5). To the mixture were added 10 µl of Ms anti B red cells surface antigen (20 mg/ml) and 1 mg of 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide. The mixture was rotated end to end at room temperature for two hours. The particles were separated magnetically, washed once with phosphate buffer and resuspended in 2 ml of phosphate buffered saline (0.1 M, pH 7.0) to give a 2.5 % w/v suspension.

Example 32

Same procedures as described in Example 31 were followed except using 40 µl of Ms anti A red cells surface antigen (5 mg/ml) to give 2 ml of 2.5 % w/v suspension.

Blood Typing Using Magnetic Particles

Example 33

In a 5mm x 65mm test tube labeled A was placed 25 µl of 2.5 % w/v Ms anti A coated magnetic particles prepared as described in Example 32. To the other test tube labeled B was placed 25 µl of 2.5 % w/v Ms anti B coated magnetic particles prepared as described in Example 31. To both test tubes was added 50 µl of 1 % packed red blood cells prepared by 1 to 100 dilution of packed red blood cells in isotonic buffered saline. The test tubes were shaken by finger tapping for several times and placed on the top of a magnet. The results were summarized as follows:

	BLOOD TYPE			
	A	B	O	AB
TUBE A	+	-	-	+
TUBE B	-	+	-	+

Where + represent a positive reaction, meaning the red cells were agglutinated by the corresponding antibody coated magnetic particles as a result the supernatant in the test tube was clear after magnetic separation. On the other hand the supernatant of a negative reaction would remain cloudy after magnetic separation due to the absence of agglutination between the red cells and the antibody coated magnetic particles.

Immunoassays Using Magnetic Particles

Example 34

In a 2 ml microcentrifuge tube was placed 1 ml of 6 % w/v, 3 micron carboxyl magnetic particles. The particles were centrifuged for 3 minutes at 10000 rpm. The supernatant was aspirated and the particles were resuspended by vortexing with 1 ml of 5 to 100 ug/ml recombinant HBcAg in acetate buffer. The tube was rotated at room temperature for two hours and centrifuged as described before. The supernatant was aspirated and the particles were resuspended in 1 ml of overcoat solution containing acetate buffer and 2 to 10 % of normal animal serum. The tube was rotated at room temperature for 2 to 16 hours and centrifuged as described before. The supernatant was aspirated and the particles were washed three times with 1 ml of isotonic buffered saline (IBS) by centrifugation and resuspension. Finally, the particles were resuspended with 1 ml of IBS and stored at 2 to 8°C.

Example 35

To the first two columns of a 96-well microtiter plate was placed 20 ul of 0.25% w/v hepatitis B core antigen (HBcAg) coated magnetic particles prepared as described in Example 34. Sample preparation consisted of various dilutions of a HBcAb positive serum into a negative plasma, followed by a 1:100 dilution of each sample into specimen dilution buffer (SDB). The SDB contained phosphate buffer, protein stabilizers, detergent, and antimicrobial agents. To the wells containing the particles were added 50 ul of each final sample dilution. After 30 minutes incubation at 37°C, the particles were separated for two minutes on a magnetic separator and washed three times with 200 ul wash buffer containing salts and detergent. To each well containing the particles was added 50 ul of goat antihuman IgG-B-D-galactosidase conjugate (0.5 ug/ml) in diluent containing salts, protein stabilizers, glycerol, detergent and antimicrobial agents. After 15 minutes incubation at 37°C the particles were separated and washed three times as described above and resuspended in 30 ul of IBS. The particles were transferred to the first two columns of a black microtiter plate (Dynatech). To each well containing the particles was added 100 ul of a solution containing 4-methylumbelliferyl-B-galactopyranoside (MUG, Sigma). The plate was incubated at 37°C and the fluorescence intensity was measured by using a Fluorescence Concentration Analyzer (FCA, Pandex) equipped with 365 nm excitation and 450 nm emission filters at five minutes interval and 10 X gain setting. The increase in fluorescence intensity in a five minutes interval was recorded in arbitrary fluorescence unit (AFU) and presented in Table I.

TABLE I

Dilution of Positive Specimen	AFU (5 Minutes) Average of Two Wells
1:100	22687
1:1000	5933
1:5000	1516
1:8000	835
1:10000	639
1:15000	495
1:20000	427
1:25000	307

Example 36

The coupling of mouse antiHBsAg to carboxyl magnetic particles was similar to Example 30.

To the wells of a black 96-well microtiter plate (Dynatech) were added 20 μ l of 0.25% w/v, 3.2 micron, mouse antiHBsAg coated carboxyl magnetic particles in duplicate. To the wells containing the magnetic particles was added 100 μ l of neat plasma containing various amounts of HBsAg or a HBsAg-negative plasma. After 30 minutes incubation at 37°C, the particles were separated for two minutes on a magnetic separator and washed once with 100 μ l of wash buffer containing salts and detergent. To each well containing the particles was added 20 μ l of mouse antiHBsAg--B-galactosidase conjugate in diluent containing salts, protein stabilizers, glycerol, detergent and antimicrobial agents. After 15 minutes incubation at 37°C, the particles were separated and washed five times as described above. To each well containing the particles was added 50 μ l of a solution containing 4-methylumbelliferyl-B-D -galactopyranoside (MUG, Sigma). The plate was incubated at 37°C and the fluorescence intensity was measured by using a Fluorescence Concentration Analyzer (FCA, Pandex) equipped with 365 nm excitation and 450 nm emission filters or five minutes interval and 10 X gain setting. The increase in fluorescence intensity in a five minutes interval was recorded in arbitrary fluorescence unit (AFU) and presented in Table II.

TABLE II

HBsAg Conc. (nano gm)	AFU (5 Minutes) Average of Two Wells
1.0	1149
0.5	455
0.25	218
0.125	118
neg.	14

Example 37

The HIV-1 antigens from HTLV-IIIb/H-9 cells (Gallo Strain) were coupled to 3.6 micron carboxyl magnetic particles by using similar procedures as described in Example 34.

To the wells of a 96-well microtiter plate were added 20 μ l of 0.25% w/v of HIV coated magnetic particles in duplicate. To the wells containing the particles were added 50 μ l of positive, borderline and negative specimens diluted 1:100 in specimen dilution buffer (SDB) containing phosphate buffer, protein stabilizers, detergent and antimicrobial agents. After 30 minutes incubation at 37°C, the particles were separated for two minutes on a magnetic separator and washed three times with 100 μ l of washed buffer containing salts and detergent. To each well containing particles was added 50 μ l of goat antihuman-B-galactosidase (approximately 0.5 μ g/ml) conjugate in diluent containing salts, protein stabilizers, glycerol, detergent and antimicrobial agents. After 15 minutes incubation at 37°C, the particles were washed four times as described above. The particles were transferred to the black microtiter plate (Dynatech). To each well containing particles was added 100 μ l of a solution containing 4-methylumbelliferyl-B-D-galactopyranoside (MUG, Sigma). The plate was incubated at 37°C and the fluorescence intensity was measured by using a Fluorescence Concentration Analyzer (FCA, Pandex) equipped with 365 nm excitation and 450 nm emission filters at five minutes intervals and 25 X gain setting. The increase in fluorescence intensity in a five minutes interval was recorded in arbitrary fluorescence unit (AFU) and presented in Table III.

TABLE III

Anti-HIV Specimens	AFU (5 Minutes) Average of Two Wells
Positive Control	9462
Borderline Specimen	527
Negative Control	86

Cell Separation Using Magnetic ParticlesExample 38

The 4.3 micron carboxyl magnetic particles prepared as described in Example 7 were washed and sonicated in phosphate buffered saline (PBS, pH 7.7), sterilized in 70% ethanol for 10 minutes, washed three times in PBS and incubated for 48 hours at 4°C with affinity-purified sheep anti-mouse immunoglobulin antibody (SAM) at 0.5 mg/ml and

a ratio of 3.3 mg antibody/100mg particles. Before use, the antibody coated magnetic particles were washed in PBS and resuspend at the desired concentration in PBS.

Human tissue culture cALLa-positive NALM-16 leukemia cells were washed and suspended in PBS. One fraction was not treated with antibody (-MoAb). The other fraction was treated with two anti-CD10 and one anti-CD9 monoclonal antibodies (+MoAb) for 30 minutes at 4°C, washed in PBS and adjusted to 3.5×10^6 cells/ml on PBS. To two tubes, one containing the antibody treated cells (+MoAb), the other containing untreated cells (-MoAb) were added SAM coated magnetic particles at a particle to starting cell ratio of 45. The tubes were rotated at 4°C for 30 minutes. The particles were separated with a magnetic separator. The supernatant was collected and centrifuged to collect the remaining cells. The pellet was resuspended in 100 μ l of trypan blue and total cell count was made. The results were presented in Table IV.

TABLE IV

Particle/cell Ratio	Cells +/- MoAb	Cells Received	% Depletion
0	+	7.62×10^5	(Control)
45	+	2.89×10^4	96.2
45	-	7.33×10^5	4.6

Example 39

A mixture containing 576 ml of deionized water, 9 ml of styrene, and 288 ml of 3.0 % w/v magnetic metal oxide prepared as described in Example 1 was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a $65 \pm 4^\circ$ C oven for one hour. To the mixture were added 18 g of potassium persulfate and 712 ml of 5% w/v, 4.0 micron fluorescent Nile Red polystyrene particles. The bottle was resealed, evacuated and rotated for one hour, and added 45 ml of 2.0% sodium dodecylsulfate. After five more hours, 9 ml of styrene and 9 g of potassium persulfate were added to the mixture. The mixture was rotated for another fifteen hours, filtered through two layers of cheese cloth, separated magnetically, and washed several times with deionized water until the supernatant was clear. The resulting fluorescent magnetic particles were resuspended to 1580 ml with deionized water to give 3.0% w/v suspension with about 11.0% magnetic metal oxide content and 4.4 micron average size.

Example 40

The fluorescent Nile Red magnetic particles, 1.580 l of 3.0% w/v, prepared as described in Example 39, were carboxylated by adding 1.23 g of sodium dodecylsulfate, 17.50 g of potassium persulfate, and solution containing 1.2 ml of undecylenic acid and 0.024 ml of divinyl benzene in 4.8 ml of methanol. The mixture was placed in a sealed bottle, evacuated and rotated at about 60 rpm in a $55^\circ - 65^\circ$ oven for five hours. The resulting fluorescent Nile Red carboxyl magnetic particles were separated magnetically and washed several times with deionized water until the supernatant was clear. The fluorescent Nile Red carboxyl magnetic particles were resuspended to 850 ml with deionized water to give a 5.0% w/v suspension with about 11.0% magnetic metal oxide content and 4.4 micron average size.

Example 41

A mixture containing 12.4 ml of 11.28% w/v, 2.24 micron polystyrene particles, 65 ml of 2.78% w/v metal oxide prepared as described in Example 5, 75 ml of deionized water and a solution containing 0.18 g of benzoyl peroxide, 7 mg of Nile Red, and 0.75 ml of divinyl benzene in 6.75 ml of styrene was placed in a sealed bottle. The bottle was evacuated and rotated at about 60 rpm in a $60^\circ - 70^\circ$ oven for about fifteen hours. The mixture was filtered through two layers of cheese cloth, separated magnetically, and washed several times with deionized water until the supernatant was clear. The resulting fluorescent crosslinked magnetic particles were resuspended to 170 ml with deionized water to give 5.4% w/v suspension with about 16.5% w/v metal oxide content and 4.0 micron average size.

Example 42

The coupling of Goat anti HBsAg to fluorescent and nonfluorescent carboxyl magnetic particles (having about the same as metal oxide content) was similar to Example 30.

To the wells of black 96-well microtiter plate (Pandex™) were added 20 μ l of 0.125% w/v, 4.0 micron, Goat anti HBsAg coated thorough mixed fluorescent and nonfluorescent carboxyl magnetic particles (at the ratio 1:1). To the wells containing the magnetic particles was added 100 μ l of neat plasma containing various amounts of HBsAg or

HBsAg-negative plasma. After thirty minutes of incubation at 37°C, the particles were separated on a magnetic separator and washed twice with 100 µl of wash buffer. To each well containing particles was added 20 µl of mouse anti-HBsAg conjugated to B-galactosidase in dilution buffer. After fifteen minutes incubation at 37°C, the particles were separated and washed six times as described above. To each well containing particles was added 50 µl of a solution containing 4-methylumbelliferyl-B-D-galactopyranoside (MUG, Sigma). The plate was incubated at 37°C and fluorescence intensity was measured by using a Fluorescence Concentration Analyzer (FCA, Pandex™) equipped with 525 nm excitation and 580 nm emission filters (Channel C, reference Channel) at eight minutes interval and 25 X gain setting. The fluorescence intensity in Channel C was recorded in arbitrary fluorescence (AFU) and presented in Table 1. The results showed that the fluorescent magnetic particles can detect the empty wells and also indicate the wells with less than average fluorescent intensity; due to the pipetting error or particle loss during the assay.

Table I

Nos. of Well	Channel C (Reference Channel)	
	Δ AFU Range	Δ AFU Range
12 Empty	4832 - 4900	4867
17	29826 - 33200	31480
1	---	19458*
1	---	27952

* AFU 19458 for one well compare to average AFU 31480 for 17 wells indicative of either loss of particles from particular well or less number of particles delivered at the start of the assay.

Example 43

Same procedures as described in Example 42 were followed except fluorescent and nonfluorescent carboxylated magnetic particles coated with Goat anti HBsAg, used simultaneously in the assay to compare the assay performance. The fluorescent intensity was measured by using Channel D (assay channel 365 nm excitation and 450 nm emission filters) and presented in Table II. The results showed that both fluorescent and nonfluorescent particles performed equally in the assay.

Table II

HBsAg Conc.	AFU	
	Fluorescent Particles	Nonfluorescent Particles
High	27238	30059
Medium	5820	5976
Low	1688	1816
Negative	326	403

Claims

1. A process to determine the presence or concentration of an analyte comprising:

- a) contacting fluorescent magnetic particles having a ligand specific for said analyte attached to said fluorescent magnetic particle with fluid specimen to form a suspension;
- b) incubating said suspension until sufficient analyte has reacted with said specific ligand;
- c) separating said magnetic particles from said suspension;
- d) adding a second labelled ligand specific for said analyte to said separated magnetic particles;
- e) incubating said suspension until sufficient analyte has reacted with said second labelled ligand specific for said analyte;
- f) separating said magnetic particles from said suspension;
- g) detecting or measuring duplex formation on said magnetic particles by means of said label; and
- h) relating the amount of labelled ligand measured with the amount of analyte measured for a control sample, wherein said fluorescent magnetic particles are used to monitor the number of particles present during said process.

2. A process to determine the presence or concentration of specific nucleic acid sequences in nucleic acid target molecules comprising:
 - a) contacting fluorescent magnetic particles having a nucleic acid complementary to said nucleic acid sequence of said target molecule attached to said fluorescent magnetic particle with fluid specimen to form a suspension;
 - b) incubating said suspension under hybridization conditions for a period of time sufficient to permit hybridization;
 - c) separating said fluorescent magnetic particles from said suspension;
 - d) adding a second labelled nucleic acid sequence complementary to said nucleic acid sequence of said target molecule;
 - e) incubating said suspension under hybridizing conditions for a period of time sufficient to permit hybridization;
 - f) separating said fluorescent magnetic particles from said suspension; and
 - g) detecting or measuring duplex formation on said magnetic particles by means of said label, wherein said fluorescent magnetic particles are used to monitor the number of particles present during said process.
3. The process of Claim 1 or 2, wherein the fluorescent material is selected from Nile™ red, Coumarin™6, Coumarin™4, Rhodamine™B, Nile™ blue, oxazine 725, oxazine 750, or a mixture of two or more of these fluorescent materials.
4. The process of Claim 1, 2 or 3, wherein the number of said particles is monitored by measuring the fluorescence intensity of fluorescent particles prior to contacting said particles with said fluid specimen and by measuring the fluorescent intensity of fluorescent particles after measuring the amount of labelled ligand associated with said fluorescent magnetic particles.
5. The process of any preceding claim wherein the magnetic particles comprise monodispersed fluorescent magnetic particles of a uniform size distribution and magnetic content comprising:
 - a) an inner fluorescent core polymer particle able to adsorb a monomer and a magnetically responsive metal oxide and polymer combination said polymer being comprised of monomers able to adsorb to said inner core polymer particle;
 - b) said metal oxide and polymer combination evenly coating said inner core particle;
 - c) said magnetic particles being of a uniform size distribution, uniform magnetic content, and monodispersed in solution, and
 - d) including an outer polymer coating said magnetically responsive metal oxide and polymer combination and a layer of functionalized polymer covering said outer polymer coating.
6. The process of Claim 5 or 6 wherein the polymer of said magnetically responsive metal oxide and polymer combination is selected from polystyrene, cross-linked polystyrene and functionalized polystyrene.
7. The process of any of claims 5-7 wherein said functionalized polymer is selected from the group of compounds which provide carboxyl, amino or hydroxyl functional groups for coupling to biological material.
8. The process of any of claims 5-8 wherein said polymer of said metal oxide and polymer combination contains a fluorescent dye or combination of fluorescent dyes.

Patentansprüche

1. Verfahren zur Bestimmung des Vorhandenseins oder der Konzentration eines Analyten, welches folgende Verfahrensschritte aufweist:
 - a) Kontaktieren fluoreszierender magnetischer Teilchen, welche einen für den Analyten spezifischen Liganden, der an die fluoreszierenden magnetischen Teilchen gebunden ist, aufweisen, mit einer flüssigen Probe zur Bildung einer Suspension;
 - b) Inkubieren der Suspension, bis eine ausreichende Menge des Analyten mit dem spezifischen Liganden

umgesetzt ist;

c) Abscheiden der magnetischen Teilchen aus der Suspension;

d) Zusetzen eines markierten zweiten Liganden, der für den Analyten spezifisch ist, zu den abgeschiedenen magnetischen Teilchen;

e) Inkubieren der Suspension, bis eine ausreichende Menge des Analyten mit dem markierten zweiten, für den Analyten spezifischen Liganden umgesetzt ist;

f) Abscheiden der magnetischen Teilchen aus der Suspension;

g) Erfassen bzw. Messen der doppelten Komplexbildung an den magnetischen Teilchen mit Hilfe der Markierung; und

h) Herstellen einer Beziehung zwischen der gemessenen Menge des markierten Liganden mit der an einer Referenzprobe gemessenen Menge des Analyten, wobei die fluoreszierenden magnetischen Teilchen zur Kontrolle der während des Verfahrens vorhandenen Anzahl von Teilchen herangezogen werden.

2. Verfahren zur Bestimmung des Vorhandenseins oder der Konzentration spezieller Nukleinsäuresequenzen in Nukleinsäure-Targetmolekülen, welches folgende Verfahrensschritte aufweist:

a) Kontaktieren fluoreszierender magnetischer Teilchen, welche eine zu der Nukleinsäuresequenz des an die fluoreszierenden magnetischen Teilchen gebundenen Targetmoleküls komplementäre Nukleinsäure aufweisen, mit einer flüssigen Probe zur Bildung einer Suspension;

b) Inkubieren der Suspension unter Hybridisierungsbedingungen über eine ausreichend lange Zeit, um eine Hybridisierung zu ermöglichen;

c) Abscheiden der magnetischen Teilchen aus der Suspension;

d) Zusetzen einer markierten zweiten Nukleinsäuresequenz, die zu der Nukleinsäuresequenz des Targetmoleküls komplementär ist;

e) Inkubieren der Suspension unter Hybridisierungsbedingungen über eine ausreichend lange Zeit, um eine Hybridisierung zu ermöglichen;

f) Abscheiden der magnetischen Teilchen aus der Suspension; und

g) Erfassen bzw. Messen der doppelten Komplexbildung an den magnetischen Teilchen mit Hilfe der Markierung, wobei die fluoreszierenden magnetischen Teilchen zur Kontrolle der während des Verfahrens vorhandenen Anzahl von Teilchen herangezogen werden.

3. Verfahren nach Anspruch 1 oder 2, bei welchem das fluoreszierende Material aus der Gruppe Nile™-Rot, Coumarin™6, Coumarin™4, Rhodamin™B, Nile™-Blau, Oxazin 725, Oxazin 750 gewählt ist oder ein Gemisch von zwei oder mehr dieser fluoreszierenden Stoffe ist.

4. Verfahren nach Anspruch 1, 2 oder 3, bei welchem die Anzahl der Teilchen durch Messen der Fluoreszenzintensität der fluoreszierenden Teilchen kontrolliert wird, ehe die Teilchen mit der flüssigen Probe in Kontakt gebracht werden, sowie durch Messen der Fluoreszenzintensität der fluoreszierenden Teilchen nach Messung der Menge des markierten Liganden, der dem fluoreszierenden magnetischen Teilchen zugeordnet ist.

5. Verfahren nach einem der vorhergehenden Ansprüche, bei welchem die magnetischen Teilchen monodispergierte fluoreszierende magnetische Teilchen in gleichmäßiger Größenverteilung und mit gleichmäßiger Magnetisierung umfassen, bei welchem:

a) ein fluoreszierendes Innenkern-Polymerteilchen vorgesehen ist, das zur Adsorption eines Monomeren in der Lage ist, und eine Kombination aus magnetisch reagierendem Metalloxid und einem Polymer, wobei das

Polymer aus Monomeren besteht, die zur Adsorption an das Innenkern-Polymerteilchen geeignet sind;

b) wobei die Kombination aus Metalloxid und Polymer das Innenkerneteilchen gleichmäßig umhüllt;

c) wobei die magnetischen Teilchen eine gleichmäßige Größenverteilung und eine gleichmäßige Magnetisierung aufweisen und in der Lösung monodispersiert sind, und

d) ein äußeres Polymer vorgesehen ist, welches die Kombination aus magnetisch reagierendem Metalloxid und Polymer umhüllt, sowie eine Schicht aus funktionalisiertem Polymer, welches die äußere Polymerumhüllung bedeckt.

6. Verfahren nach Anspruch 5 oder 6, bei welchem das Polymer der Kombination aus magnetisch reagierendem metalloxid und Polymer aus der Gruppe Polystyrol, vernetztem Polystyrol und funktionalisiertem Polystyrol gewählt wird.

7. Verfahren nach einem der Ansprüche 5 bis 7, bei welchem das funktionalisierte Polymer aus der Gruppe von Verbindungen gewählt ist, bei welchen Carboxyl-, Amino- oder Hydroxyl-Funktionsgruppen zur Ankopplung an biologisches Material vorgesehen sind.

8. Verfahren nach einem der Ansprüche 5 bis 8, bei welchem das Polymer in der Kombination aus Metalloxid und Polymer einen fluoreszierenden Färbemittel oder eine Kombination aus fluoreszierenden Färbemitteln enthält.

Revendications

1. Procédé pour déterminer la présence ou la concentration d'un analyte, comprenant les étapes consistant à:

- a) mettre en contact des particules magnétiques fluorescentes, possédant un ligand spécifique audit analyte fixé à ladite particule magnétique fluorescente, avec un échantillon de fluide pour former une suspension;
- b) incuber ladite suspension jusqu'à ce que suffisamment d'analyte ait réagi avec ledit ligand spécifique;
- c) séparer lesdites particules magnétiques de ladite suspension;
- d) ajouter auxdites particules magnétiques séparées un second ligand marqué spécifique audit analyte;
- e) incuber ladite suspension jusqu'à ce que suffisamment d'analyte ait réagi avec ledit second ligand marqué spécifique audit analyte;
- f) séparer lesdites particules magnétiques de ladite suspension;
- g) détecter ou mesurer la formation d'un duplex sur lesdites particules magnétiques à l'aide dudit marqueur; et
- h) établir la relation entre la quantité de ligand marqué mesurée et la quantité d'analyte mesurée pour un échantillon témoin, lesdites particules magnétiques fluorescentes étant utilisées pour contrôler le nombre de particules présentes durant ledit procédé.

2. Procédé pour déterminer la présence ou la concentration de séquences d'acides nucléiques spécifiques dans des molécules cibles d'acide nucléique, comprenant les étapes consistant à:

- a) mettre en contact des particules magnétiques fluorescentes, possédant un acide nucléique complémentaire de ladite séquence d'acides nucléiques de ladite molécule cible fixé à ladite particule magnétique fluorescente, avec un échantillon de fluide pour former une suspension;
- b) incuber ladite suspension dans des conditions d'hybridation pendant une durée suffisante pour permettre l'hybridation;
- c) séparer lesdites particules magnétiques fluorescentes de ladite suspension;
- d) ajouter une seconde séquence d'acides nucléiques marquée complémentaire de ladite séquence d'acides nucléiques de ladite molécule cible;
- e) incuber ladite suspension dans des conditions d'hybridation pendant une durée suffisante pour permettre l'hybridation;
- f) séparer lesdites particules magnétiques fluorescentes de ladite suspension; et
- g) détecter ou mesurer la formation d'un duplex sur lesdites particules magnétiques à l'aide dudit marqueur, lesdites particules magnétiques fluorescentes étant utilisées pour contrôler le nombre de particules présentes durant ledit procédé.

3. Procédé selon la revendication 1 ou 2, dans lequel la substance fluorescente est choisie parmi le rouge Nile™, la Coumarine™6, la Coumarine™4, la Rhodamine™B, le bleu Nile™, l'oxazine 725, l'oxazine 750, ou un mélange d'au moins deux de ces substances fluorescentes.

4. Procédé selon la revendication 1, 2 ou 3, dans lequel le nombre desdites particules est contrôlé par la mesure de l'intensité de fluorescence des particules fluorescentes avant la mise en contact desdites particules avec ledit échantillon de fluide et par la mesure de l'intensité de fluorescence des particules fluorescentes après la mesure de la quantité de ligand marqué associé auxdites particules magnétiques fluorescentes.

5. Procédé selon l'une quelconque des revendications précédentes, dans lequel les particules magnétiques comprennent des particules magnétiques fluorescentes monodispersées, de distribution granulométrique uniforme et d'aimantation uniforme, comprenant:

- a) une particule de polymère fluorescente interne formant le noyau, susceptible d'adsorber un monomère et une combinaison d'oxyde métallique et de polymère, à sensibilité magnétique, ledit polymère étant composé de monomères susceptibles de s'adsorber sur ladite particule de polymère interne formant le noyau;
- b) ladite combinaison d'oxyde métallique et de polymère enrobant uniformément ladite particule interne formant le noyau;
- c) lesdites particules magnétiques ayant une distribution granulométrique uniforme, une aimantation uniforme, et étant monodispersées dans la solution; et
- d) renfermant un enrobage de polymère externe, ladite combinaison d'oxyde métallique et de polymère, à sensibilité magnétique, et une couche de polymère fonctionnalisé recouvrant ledit enrobage de polymère externe.

6. Procédé selon la revendication 5 ou 6, dans lequel le polymère de ladite combinaison d'oxyde métallique et de polymère, à sensibilité magnétique, est choisi parmi le polystyrène, le polystyrène réticulé et le polystyrène fonctionnalisé.

7. Procédé selon l'une quelconque des revendications 5-7, dans lequel ledit polymère fonctionnalisé est choisi dans le groupe de composés apportant des groupes fonctionnels carboxyle, amino ou hydroxyle, pour le couplage à une substance biologique.

8. Procédé selon l'une quelconque des revendications 5-8, dans lequel ledit polymère de ladite combinaison d'oxyde métallique et de polymère contient un colorant fluorescent ou une combinaison de colorants fluorescents.

FIG. 1

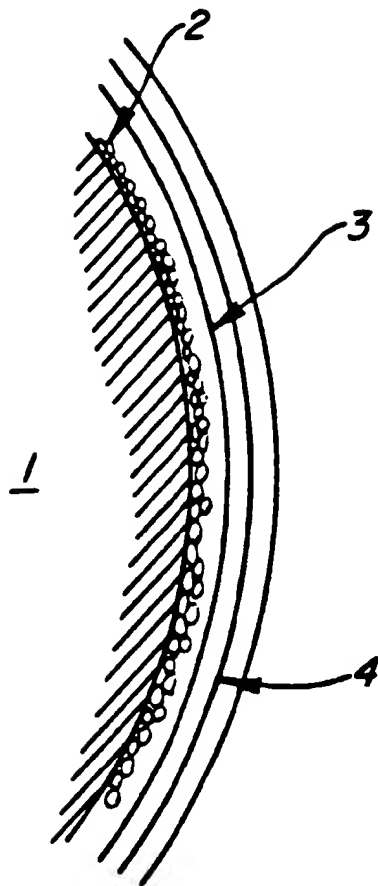




FIG. 2

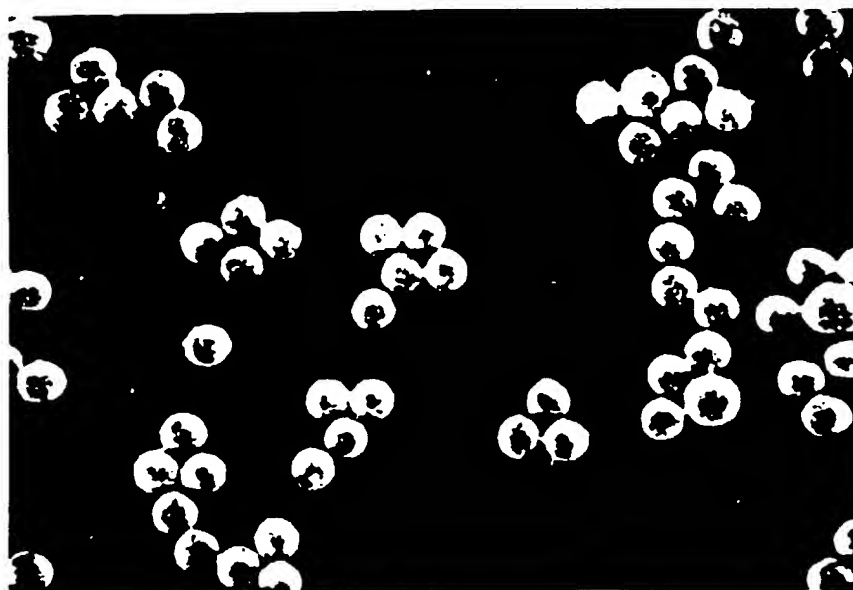


FIG. 3a

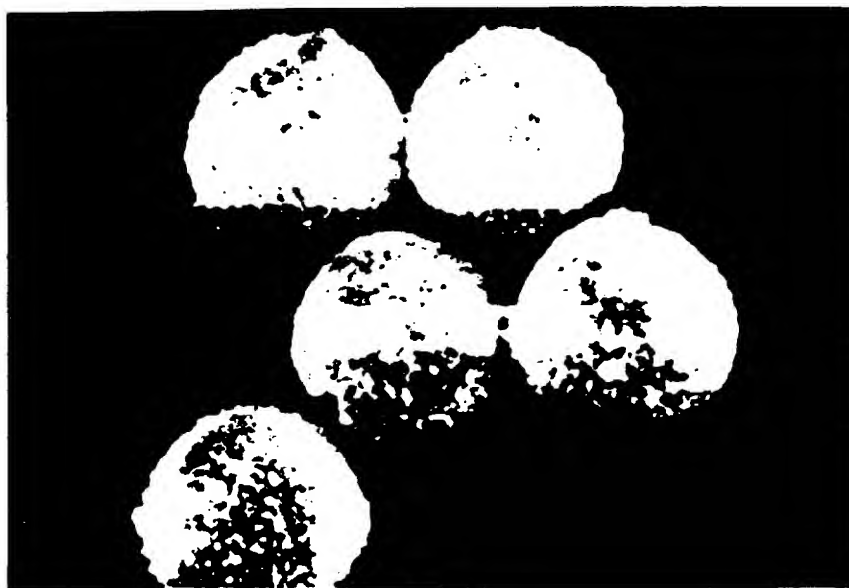


FIG. 3b